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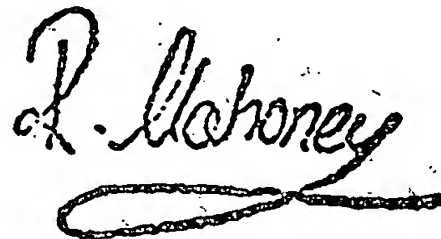

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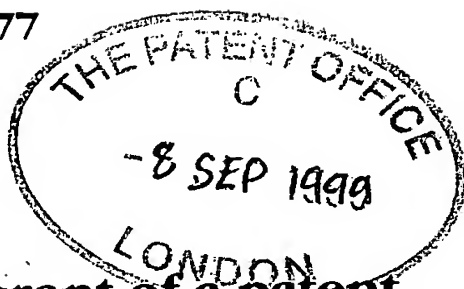
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Request for grant of a patent

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08 SEP 1999

The Patent Office

Cardiff Road
Newport
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1. Your reference

PH99052

2. Patent application number

(The Patent Office will fill in this part)

9921220.1

3. Full name, address and postcode of the or of each applicant (underline all surnames)

Rhone-Poulenc Agriculture Ltd
Fyfield Road
Ongar, Essex
CM5 0HW
ENGLAND

Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its incorporation

5623206001

4. Title of the invention

NEW HERBICIDAL COMPOSTIONS

5. Name of your agent (if you have one)

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

Rachel M Colegate Hayward
Patent Department
Rhone-Poulenc Agriculture Ltd
Fyfield Road
Ongar, Essex
CM5 0HW

Patents ADP number (if you know it)

5623206002

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number

Country

Priority application number
(if you know it)

Date of filing
(day / month / year)

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application

Date of filing
(day / month / year)

8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:

NO

- a) any applicant named in part 3 is not an inventor, or
 - b) there is an inventor who is not named as an applicant, or
 - c) any named applicant is a corporate body.
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Patents Form 1/77

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Continuation sheets of this form	-
Description	19
Claim(s)	8
Abstract	1
Drawing(s)	-



10. If you are also filing any of the following, state how many against each item.

Priority documents	-
Translations of priority documents	-
Statement of inventorship and right to grant of a patent (Patents Form 7/77)	-
Request for preliminary examination and search (Patents Form 9/77)	-
Request for substantive examination (Patents Form 10/77)	-
Any other documents (please specify)	-

11. I/We request the grant of a patent on the basis of this application.

Signature Rachel M Colegate Hayward Date 08/09/99

12. Name and daytime telephone number of person to contact in the United Kingdom
- Rachel M Colegate Hayward
01277 301201

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PH99052

PATENTS ACT 1977

SPECIFICATION

BRITISH PATENT APPLICATION ENTITLED

NEW HERBICIDAL COMPOSITIONS

in the name of Rhône-Poulenc Agriculture Ltd, Fyfield Road, Ongar, Essex.

CM5 0HW. England

New Herbicidal Compositions

Background of the Invention

The present invention relates to the safening of herbicidal compounds, in particular the safening of benzoylisoxazole and/or dione derivatives which are useful for the growing of crops in particular for maize (*Zea mays*) and to compositions useful for such treatment.

Discussion of Related Art

An important factor influencing the usefulness of a given herbicide is its selectivity toward crops. In some cases, a beneficial crop is susceptible to the effects of a herbicide when applied at application rates needed to control weed growth. In addition, certain herbicidal compounds are phytotoxic to some weed species but not to others. This may render such herbicides unsuitable for controlling weeds in the presence of certain crops. To be effective, a herbicide must cause minimal damage (preferably no damage) to the beneficial crop while maximising the damage to weed species which infest the locus of the crop. Reduction in herbicidal injury to crops without an unacceptable reduction in the herbicidal action can be accomplished by the use of crop protectants known as "antidotes" or "safeners".

Identification of an antidote which safens a herbicide in crops is a complicated task. The precise mechanism by which an antidote reduces herbicidal crop injury has not been established. In general, the safening effect of a compound is specific to the herbicidal partner and the crop where the active ingredients are applied.

Benzoylisoxazoles are known to possess herbicidal properties for example, European Patent Publication Nos. 0418175, 0487357, 0527036 and 0560482. European Patent Publication Nos. 0496630, 0496631, 0625505 and 0625508 disclose certain dione derivatives possessing herbicidal properties. In general such herbicides are very active against broad-leaved and grass weeds by pre- and/or post-emergence application. The method of controlling vegetation with these compounds comprises

applying a herbicidally effective amount of the compounds, usually with an inert carrier or diluent, to the area where herbicidal control is desired. However, the herbicidal benzoylisoxazole and/or dione compounds have been found in some instances to adversely affect or interfere with the development of crop plants, especially maize crops. The effective use of these herbicides for controlling weeds in the presence of such crops may be enhanced by, or may require in certain instances, the addition of a compound which is antidotally effective with the herbicide.

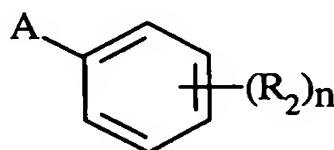
Although it is possible to say in general terms that herbicides may be used in the presence of a safener, the problem in identifying specific safeners for specific crops at appropriate rates to control weed growth, is substantial.

The applicants have found that certain compounds are effective antidotes for the protection of crops, especially maize crops, from herbicidal injury or the reduction of herbicidal injury caused by the application of an amount of a benzoylisoxazole and/or dione compound (optionally in admixture with a partner herbicide) effective to control the growth of weeds.

It is an object of the present invention to provide compositions of benzoylisoxazoles and/or dione herbicides in combination with antidotes, said compositions providing a reduction in crop injury, especially to maize (*Zea mays*), arising from the phytotoxicity of the herbicides.

Description of the Invention

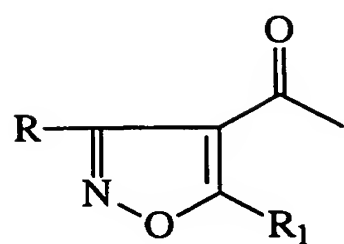
The present invention provides a method of reducing phytotoxicity to a crop (especially maize) at a locus caused by the application thereto of a herbicidal benzoylisoxazole and/or dione derivative of formula (I):



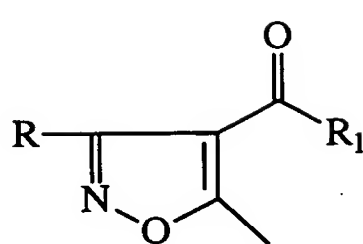
(I)

wherein:

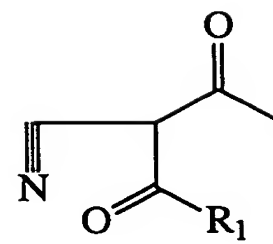
A represents a group (A-1) to (A-7):



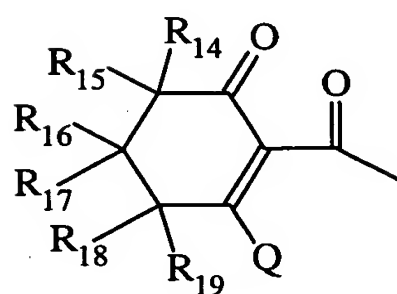
(A-1)



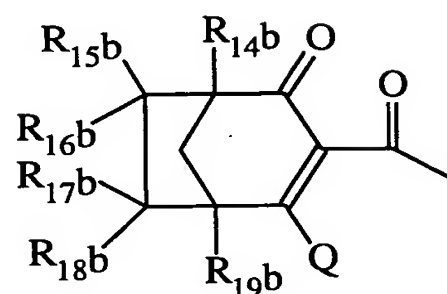
(A-2)



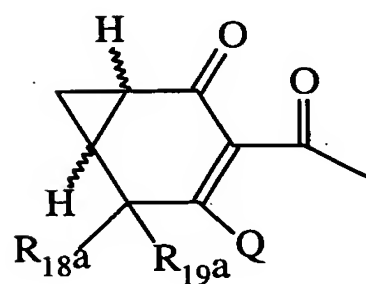
(A-3)



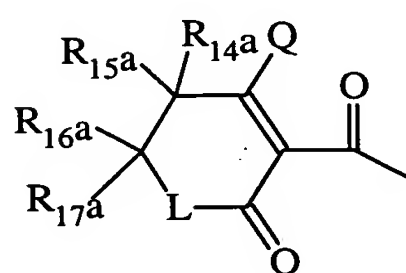
(A-4)



(A-5)

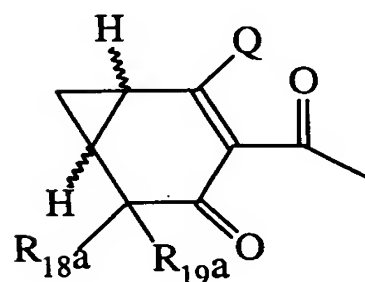


(A-6)

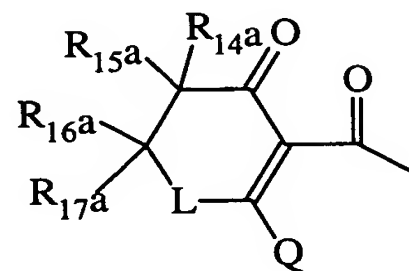


(A-7)

or a corresponding formula (A-6a) or (A-7a):



(A-6a)



(A-7a)

in which the position of the carbonyl group and the group Q are reversed and the double bond in the ring is attached to the carbon atom attached to the group Q;

5 R represents a hydrogen atom or a halogen atom; a straight- or branched chain alkyl, alkenyl or alkynyl group containing from one to six carbon atoms which is optionally substituted by one or more halogen atoms; a cycloalkyl group containing from 3 to 6 carbon atoms optionally substituted by one or more groups R^5 , one or more halogen atoms or a group $-CO_2R^3$; or a group selected from $-CO_2R^3$, $-COR^5$,
10 cyano, nitro, $-CONR^3R^4$ and $-S(O)_kR^{13}$;

R^1 represents a straight- or branched-chain alkyl, alkenyl or alkynyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms; or a cycloalkyl group containing from three to six carbon atoms optionally substituted by one
15 or more groups R^5 or one or more halogen atoms;

R^2 represents a halogen atom; a straight- or branched-chain alkyl, alkenyl or alkynyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms; a straight- or branched-chain alkyl group containing up to six carbon atoms which is
20 substituted by one or more groups $-OR^5$; or a group selected from nitro, cyano, $-CO_2R^5$, $-S(O)_pR^6$, $-O(CH_2)_mOR^5$, $-COR^5$, $-NR^{11}R^{12}$, $-N(R^8)SO_2R^7$, $-N(R^8)CO_2R^7$, $-OR^5$, $-OSO_2R^7$, $-SO_2NR^3R^4$, $-CONR^3R^4$, $-CSNR^3R^4$, $-(CR^9R^{10})_t-S(O)_qR^7$ and $-SF_5$;

25 or two groups R^2 , on adjacent carbon atoms of the phenyl ring may, together with the carbon atoms to which they are attached, form a 5 to 7 membered saturated or unsaturated heterocyclic ring containing up to three ring heteroatoms selected from nitrogen, oxygen and sulfur, which ring is optionally substituted by one or more groups selected from halogen, nitro, $-S(O)_pR^{13}$, C_{1-4} alkyl, C_{1-4} alkoxy, C_{1-4} haloalkyl,

C₁₋₄ haloalkoxy, =O (or a 5- or 6- membered cyclic acetal thereof), and =NO-R³, it being understood that a sulphur atom, where present in the ring, may be in the form of a group -SO- or -SO₂-;

n represents an integer from one to five: when n is greater than one the groups R² may be the same or different;

R³, R⁴ and R²² each independently represent a hydrogen atom, or a straight- or branched chain alkyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms;

R⁵ and R²³ each independently represent a straight- or branched-chain alkyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms or a straight- or branched-chain alkenyl or alkynyl group containing from two to six (preferably from three to six) carbon atoms which is optionally substituted by one or more halogen atoms;

R⁶ and R⁷, which may be the same or different, each represent R⁵; or phenyl optionally substituted by from one to five groups which may be the same or different selected from a halogen atom, a straight- or branched-chain alkyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms, nitro, cyano, -CO₂R⁵, -S(O)_pR¹³, -NR¹¹NR¹², -OR⁵ and -CONR³R⁴;

R⁸, R⁹ and R¹⁰ each represent a hydrogen atom or R⁶;

R¹¹ and R¹² each represent hydrogen or R⁵;

R¹³ and R²¹ represent a straight- or branched-chain alkyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms;

Q represents hydroxy, C1-6 alkoxy, OR²⁰, SR²⁰ or SR²¹;

L represents oxygen or NR²²;

R¹⁴, R^{14a}, R^{14b}, R¹⁵, R^{15a}, R^{15b}, R¹⁶, R^{16a}, R^{16b}, R¹⁷, R^{17a}, R^{17b}, R¹⁸, R^{18a}, R^{18b}, R¹⁹, R^{19a} and R^{19b} represent the same

or different groups selected from hydrogen, R^{23} , $-(CH_2)_uCO_2R^{22}$,
halogen, cyano, C1-6 alkoxy, $-(CH_2)_x$ -[phenyl optionally substituted by
from one to five groups R^{24} which may be the same or different], and
cycloalkyl containing from three to six carbon atoms optionally
5 substituted by C1-6 alkyl or $-S(O)_pR^{21}$;

R^{20} represents phenyl optionally substituted by from one to five
groups selected from halogen, C1-6 alkyl, C1-6 haloalkyl, C1-6 alkoxy
and nitro;

10 R^{24} represents a group selected from halogen, R^{25} , nitro, cyano,
 $-CO_2R^{26}$, $-S(O)_pR^{21}$, $-OR^{21}$ and $-NR^{26}R^{27}$;

R^{25} represents a straight- or branched- chain alkyl group
containing one to three carbon atoms optionally substituted by one or
more halogen atoms;

15 R^{26} and R^{27} which may be the same or different, each represents
hydrogen or R^{23} ;

p, q and u independently represent the values zero, one or two;

k and m represent one, two or three;

x represents zero or one;

20 t represents an integer from one to four; when t is greater than one,
the groups R^9 and R^{10} may be the same or different;

or an agriculturally acceptable salt or metal complex thereof;
which method comprises applying to the locus of the crop an antidotally
effective amount of an antidote compound, optionally with a partner
herbicide.

25 It will be understood that the said antidote is, in general,
antidotally effective for said benzoylisoxazole and/or dione derivative.

It will be understood that antidotes used in the method of the
invention may form for example salts, and that the use of such salts is
also embraced by the invention.

30 In this patent specification including the accompanying claims it is
understood that the term 'agriculturally acceptable salts' is meant salts

the cations or anions of which are known and accepted in the art for the formation of salts for agricultural or horticultural use. Preferably the salts are water-soluble. Suitable salts with bases include alkali metal (e.g. sodium and potassium), alkaline earth metal (e.g. calcium and magnesium), ammonium and amine (e.g. diethanolamine, triethanolamine, octylamine, morpholine and dioctylmethylamine) salts. Suitable acid addition salts, e.g. formed by compounds of formula (I) containing an amino group, include salts with inorganic acids, for example hydrochlorides, sulphates, phosphates and nitrates and salts with organic acids for example acetic acid.

It will be understood that the term 'dione' as used in this specification including the accompanying claims does not exclude the possible presence of additional C = O groups as in triones.

Compounds of formula (I) may exist in enolic tautomeric forms that may give rise to geometric isomers around the enolic double bond. Furthermore in certain cases the above substituents may contribute to optical isomerism and/or stereoisomerism. All such forms and mixtures thereof are embraced by the present invention.

It is to be understood that in this specification compounds comprising a cyclohexane ring corresponding to formula (A-6) or (A-7) or a precursor thereof include the compounds with the corresponding formula (A-6a) or (A-7a) or precursors thereof.

In the definitions of symbols in this specification including the accompanying claims unless otherwise specified the following term is generally defined thus:-

'halogen' means a fluorine, chlorine, bromine or iodine atom.

Alkyl groups and moieties are straight or branched chain and contain from 1 to 6 carbon atoms.

Preferably A represents a group of formula (A-1), (A-2), (A-3) or (A-4) (compounds of formula (A-1) are most preferred).

The benzoyl ring of the compounds of formula (I) is preferably 2,4-disubstituted, 2,3-disubstituted or 2,3,4-trisubstituted.

Preferably in formulae (A-4) to (A-7), the groups R^{14} , R^{15} , R^{16} , R^{17} , R^{18} , R^{19} , R^{14a} , R^{15a} , R^{16a} , R^{17a} , R^{14b} , R^{15b} , R^{16b} , R^{17b} , R^{18b} and R^{19b} represent hydrogen or lower alkyl (preferably hydrogen, methyl or ethyl); L (in A-7a) represents NH; and Q represents hydroxy or -S-phenyl.

Compounds of formula (I) in which A represents (A-1), (A-2) or (A-3); R represents hydrogen or $-CO_2R^3$ (in A-1 or A-2) wherein R^3 represents or a straight- or branched chain alkyl group containing up to three carbon atoms; and R^1 represents cyclopropyl are preferred.

A further preferred class of compounds of formula (I) wherein A represents (A-1) are those wherein:

R is hydrogen or $-CO_2Et$;

R^1 is cyclopropyl;

and two groups R^2 , on adjacent carbon atoms of the phenyl ring may, together with the carbon atoms to which they are attached, combine to form a 5 or 6 membered saturated or unsaturated heterocyclic ring which is fused to the 2,3 or 3,4 positions of the benzoyl ring; wherein the heterocyclic ring contains two hetero atoms selected from sulphur and oxygen which are attached to the 2 and 3, or 3 and 4 positions of the benzoyl ring; and in which the 4-substituent of the benzoyl ring is halogen or $S(O)_pMe$, or the 2-substituent of the benzoyl ring is methyl, $S(O)_pMe$ or $-CH_2S(O)_qMe$ respectively; and optionally the heterocyclic ring may be substituted by one or more halogen atoms.

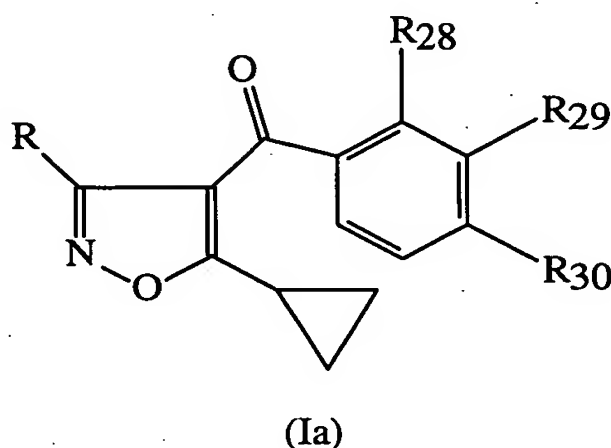
A further preferred class of compounds of formula (I) are those wherein A represents (A-1); R is hydrogen or $-CO_2Et$; R^1 is cyclopropyl; R^2 is a halogen atom or a group selected from $-CF_3$, Me, Et, $-S(O)_pMe$, $-CH_2S(O)_qMe$ and optionally halogenated methoxy or ethoxy; and n is two or three.

A further preferred class of compounds of formula (I) wherein A represents (A-4) are those wherein:

R^{14} , R^{15} , R^{16} , R^{17} , R^{18} and R^{19} represent hydrogen;

and two groups R^2 , on adjacent carbon atoms of the phenyl ring may, together with the carbon atoms to which they are attached, combine to form a 6 membered saturated heterocyclic ring which is fused to the 2,3 or 3,4 positions of the benzoyl ring; wherein the heterocyclic ring contains a sulphur atom attached to the 4 position of the benzoyl ring, optionally in the form of a group -SO- or -SO₂-, and which ring is substituted by a 5- or 6- membered acetal thereof.

A more preferred class of compounds of formula (I) having the formula (Ia):



are those wherein:

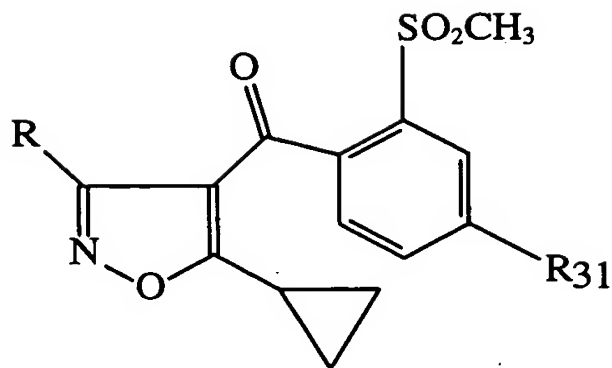
R is hydrogen or -CO₂Et;

R^{28} is selected from -S(O)_pMe, Me, Et, a chlorine, bromine or fluorine atom, methoxy, ethoxy and -CH₂S(O)_qMe;

R^{29} is selected from a hydrogen atom, a chlorine, bromine or fluorine atom, methoxy, ethoxy and -S(O)_pMe; and

R^{30} is selected from a hydrogen atom, a chlorine, bromine or fluorine atom, methoxy and CF₃.

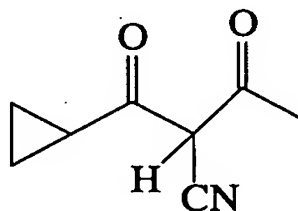
An especially preferred class of compounds of formula (I) have the formula (Ib):



(Ib)

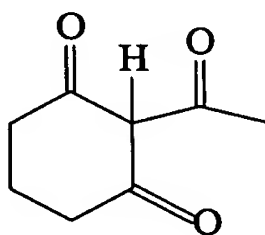
wherein R^{31} is chlorine, bromine or trifluoromethyl; and
R is hydrogen or $-CO_2Et$.

5 Preferred diones are those in which a substituted phenyl ring as defined in formula (I); (Ia); or (Ib), is attached to a grouping;



10 Such diones in which the phenyl ring is substituted by two groups independently selected from halogen, alkyl, $S(O)_t$ alkyl ($t = 0, 1$ or 2) or haloalkyl are also preferred.

Preferred triones are those in which a substituted phenyl ring, as defined above, is attached to a grouping;



15 The following compounds of formula (I) are among the most preferred for use in the present invention:

5-cyclopropyl-4-[2-chloro-3-ethoxy-4-(ethylsulphonyl)benzoyl]isoxazole;

4-(4-chloro-2-methylsulphonylbenzoyl)-5-cyclopropylisoxazole;

20 5-cyclopropyl-4-(2-methylsulphonyl-4-trifluoromethylbenzoyl)isoxazole;

4-(4-bromo-2-methylsulphonylbenzoyl)-5-cyclopropylisoxazole;

5-cyclopropyl-4-[4-fluoro-3-methoxy-2-(methylsulphonyl)benzoyl]isoxazole;

4-(4-bromo-2-methylsulphonylmethylbenzoyl)-5-cyclopropylisoxazole;

5 ethyl 5-cyclopropyl-4-(2-methylsulphonyl-4-trifluoromethylbenzoyl)isoxazole-3-carboxylate;

2-[2-chloro-(4-methylsulphonyl)benzoyl]-1,3-cyclohexanedione;

2-[2-nitro-(4-methylsulphonyl)benzoyl]-1,3-cyclohexanedione;

10 2-(2,3-dihydro-5,8-dimethyl-1,1-dioxospiro[4H-1-benzothiine-4,2'-[1,3]dioxolan]-6-ylcarbonyl)cyclohexane-1,3-dione;

5-cyclopropyl-4-(2-methylsulphonyl-4-trifluoromethylbenzoyl)-3-methylthio-isoxazole;

2-cyano-3-cyclopropyl-1-(2-methylsulphonyl-4-trifluoromethylphenyl)propan-1,3-dione.

15 The most preferred compounds are 5-cyclopropyl-4-(2-methylsulphonyl-4-trifluoromethylbenzoyl)isoxazole and 2-[2-nitro-(4-methylsulphonyl)benzoyl]-1,3-cyclohexanedione.

Herbicidal benzoylisoxazole and/or dione compounds for use in this invention may be prepared by the methods described in the
20 aforementioned patent publications, or by the application or adaptation of known methods used or described in the chemical literature.

It has been found that such antidote compounds can be selected from a wide range of chemical substances. The preferred compositions of this invention may include one or more antidotes which are not
25 suggested by earlier safeners which have been proposed for use with the benzoylisoxazole and/or dione derivatives of formula (I). The compositions of the invention may include one or more of the following safeners:

30 flurazole, which is benzyl 2-chloro-4-trifluoromethyl-1,3-thiazole-5-carboxylate; fenclorazole-ethyl, which is ethyl 1-(2,4-dichlorophenyl)-5-trichloromethyl-1*H*-1,2,4-triazole-3-carboxylate; fenchorazole, which is ethyl 1-(2,4-dichlorophenyl)-5-trichloromethyl-1*H*-1,2,4-triazole-3-carboxylic acid; benoxacor, which is (±)-4-

dichloroacetyl-3,4-dihydro-3-methyl-2*H*-1,4-benzoxazine; dichlormid, which is *N,N*-diallyl-2,2-dichloroacetamide; fenclozim, which is 4,6-dichloro-2-phenylpyrimidine; furilazole, which is (*RS*)-3-dichloroacetyl-5-(2-furyl)-2,2-dimethyloxazolidine; mefenpyr-diethyl, which is diethyl (*RS*)-1-(2,4-dichlorophenyl)-5-methyl-2-pyrazoline-3,5-dicarboxylate; CMPI, which is *N*-(4-chlorophenyl)maleimide; 4-hydroxy-1-methyl-3-(1-1*H*-tetrazol-5-ylmethanoyl)-1*H*-quinolin-2-one; daimuron, which is 1-(1-methyl-1-phenylethyl)-3-*p*-tolylurea; (*S*)-MBU, which is (*S*)-1-(1- α -methylbenzyl)-3-*p*-tolylurea; dimepiperate, which is *S*-1-methyl-1-phenylethyl piperidine-1-carbothioate; 5,5-diphenylisoxazolinone-3-carboxylic acid; and ethyl 5,5-diphenylisoxazolinone-3-carboxylate.

The mixtures of the invention may be used to obtain selective weed control with low crop injury in various crop plants such as maize, soybean, cotton, canola, sugar beet, potatoes, wheat, tobacco, rice and oil seed rape. Preferred crops include maize, sugar beet, cotton and canola. Particularly preferred crop species are maize and soybean, especially maize.

Effective weed control coupled with low crop injury is a result of treatment of a plant locus with a combination of a herbicidal benzoylisoxazole and/or dione derivative and an antidote compound in accordance with the present invention. By application to the 'plant locus' is meant application, for example to the plant growing medium, such as soil, as well as to the seeds, emerging seedlings, roots, stems, leaves or other plant parts.

The phrase 'combination of a herbicidal isoxazole and/or dione derivative and an antidote compound' includes various methods of treatment. For example, the soil of a plant locus may be treated with a "tank-mix" composition containing a mixture of the herbicide and the antidote which is "in combination", or the soil may be treated with the herbicide and antidote compounds separately so that the "combination" may be made on, or in the soil. After such treatments of the soil with a mixture of herbicide and antidote or by separate or sequential application of the herbicide and the antidote to the soil, the herbicide

and antidote may be mixed into or incorporated into the soil either by mechanical mixing of the soil with implements or by "watering in" by rainfall or irrigation. The soil of a plant locus may also be treated with antidote by application of the antidote in a dispersible-concentrate form such as a granule. The granule may be applied to a furrow which is prepared for receipt of the crop seed and the herbicide may be applied to the plant locus either before or after in-furrow placement of the antidote-containing granule so that the herbicide and antidote form a "combination". Crop seed may be treated or coated with the antidote compound either while the crop seed is in-furrow just after seeding or, more commonly, the crop seed may be treated or coated with antidote prior to seeding into a furrow. The herbicide may be applied to the soil plant locus before or after seeding and a "combination" is made when both herbicide and an antidote-coated seed are in the soil.

In one embodiment the method of the invention is preferably performed by applying the antidote directly to the seed before planting. This is generally effected by coating a quantity of crop seed with the antidote and thereafter planting the coated seed.

The amount of a particular benzoylisoxazole and/or dione herbicide to be applied to the plant locus or crop-growing area will depend upon the nature of the weeds, the particular herbicide used, the time of application, the climate and the nature of the crop. Application rates of from about 0.004kg ha^{-1} to 5kg ha^{-1} herbicide are generally suitable, with a rate of about 0.01kg ha^{-1} to 2kg ha^{-1} being preferred.

The amount of antidote used in the method of the invention varies according to a number of parameters including the particular antidote employed, the crop to be protected, the amount and rate of herbicide applied, and the edaphic and climatic conditions prevailing. Also, the selection of the specific antidotes for use in the method of the invention, the manner in which it is to be applied and the determination of the activity which is non-phytotoxic but antidotally effective, can be readily performed in accordance with common practice in the art.

The antidote is applied in combination with the herbicide in a non-phytotoxic antidotally effective amount. By "non-phytotoxic" is meant an amount of the antidote which causes at most minor or no injury to the desired crop species. By "antidotally-effective" is meant an antidote used in an amount which is effective as an antidote with the herbicide to decrease the extent of injury caused by the herbicide to the desired crop species.

The following non-limiting examples illustrate the invention wherein Safener A is ethyl 5,5-diphenylisoxazolinone-3-carboxylate and Safener B is 5,5-diphenylisoxazolinone-3-carboxylic acid.

Example 1

Maize seeds were sown in non-sterile loam and safener, dissolved in acetone was applied to the soil surface. After 30 minutes a treatment of herbicide, Compound A [5-cyclopropyl-4-(2-methylsulphonyl-4-trifluoromethylbenzoyl) isoxazole] was applied to the treated soil.

A visual assessment of the percentage phytotoxicity (measured as a reduction in green plant matter or plant height) compared to an untreated control was made 14 days after treatment (DAT).

Maize seeds were sown in non-sterile loam and grown up to a 1.5 - 2 leaf stage. Safener, dissolved in acetone, was applied post-emergence to the soil surface. After 30 minutes a treatment of herbicide, Compound A [5-cyclopropyl-4-(2-methylsulphonyl-4-trifluoromethylbenzoyl) isoxazole] was applied to the treated soil.

A visual assessment of the percentage phytotoxicity compared with an untreated control was made 14 DAT.

Pre-emergence activity of Compound A on maize in the presence of safeners.

	<u>[safener]</u> <u>g/ha</u>	<u>% phytotoxicity</u>
Cpd A	63	15
mefenpyr-diethyl	63	8.75
fenchlorazole-ethyl	63	11.3
Safener A	63	11.3
Safener B	63	8.75

Post-emergence activity of Compound A (63g/ha) on maize in the presence of safeners.

	<u>[safener]</u> <u>g/ha</u>	<u>% phytotoxicity</u>
Cpd A	63	27.5
mefenpyr-diethyl	63	25
fenchlorazole-ethyl	63	27.5
Safener A	63	5
Safener B	63	5

Post-emergence activity of Compound A (125g/ha) on maize in the presence of safeners.

	<u>g/ha</u>	<u>% phytotoxicity</u>
Cpd A	125	43
Safener A	31	15
	63	10
	125	15
Safener B	31	19
	63	18
	125	25

According to a further feature of the present invention, there are provided herbicidal compositions comprising:

(a) a herbicidally effective amount of a benzoylisoxazole and/or dione derivative of formula (I) or an agriculturally acceptable salt or metal complex thereof; and

(b) an antidotally effective amount of an antidote compound or an agriculturally acceptable salt thereof;

in association with, and preferably homogeneously dispersed in, one or more compatible herbicidally- acceptable diluents or carriers and/or surface-active agents (i.e. diluents or carriers or surface-active agents of the type generally acceptable in the art as being suitable for use in herbicidal compositions and which are compatible with the herbicides of the present invention). The term "homogeneously dispersed" is used to include compositions in which the benzoylisoxazole and/or dione of formula (I) and antidote are dissolved in the other components. The term "herbicidal composition" is used in a broad sense to include not only compositions which are ready for use as herbicides but also concentrates which must be diluted before use.

The ratio of herbicide to antidote may vary depending upon the crop to be protected, weed to be inhibited, herbicide used, etc., but normally an herbicide-to-antidote ratio ranging from 1:25 to 60:1 parts by weight may be employed, although much higher rates of antidote may be used, e.g., 1:100 to 1:300 parts by weight of herbicide to-antidote. The preferred weight ratio of herbicide-to-antidote is from 1:10 to 30:1. Another preferred weight range ratio is from 1:1 to 20:1, with an even more preferred weight ratio range from 2:1 to 15:1.

Preferably, the compositions contain from 0.05 to 90% by weight of benzoylisoxazole and/or dione of formula (I) and antidote.

The herbicidal composition may contain solid and liquid carriers and surface-active agents (e.g. wetters, dispersants or emulsifiers alone or in combination). Surface-active agents that may be present in the herbicidal compositions of the present invention may be of the ionic or non-ionic types, for example sulphuricinoleates, quaternary ammonium derivatives, products based on condensates of ethylene oxide with nonyl- or octyl-phenols, or carboxylic acid esters of anhydrosorbitols which have been rendered soluble by etherification of the free hydroxy groups by condensation with ethylene oxide, alkali and alkaline earth metal salts of sulphuric acid esters and sulphonic acids such as dinonyl- and dioctyl-sodium sulphonosuccinates and alkali and alkaline earth metal salts of high molecular weight sulphonic acid derivatives such as

sodium and calcium lignosulphonates. Examples of suitable solid diluents or carriers are aluminium silicate, talc, calcined magnesia, kieselguhr, tricalcium phosphate, powdered cork, absorbent carbon black and clays such as kaolin and bentonite. Examples of suitable liquid diluents include water, acetophenone, cyclohexanone, isophorone, toluene, xylene, and mineral, animal, and vegetable oils (these diluents may be used alone or in combination).

Herbicidal compositions according to the present invention may also contain, if desired, conventional adjuvants such as adhesives, protective colloids, thickeners, penetrating agents, stabilisers, sequestering agents, anti-caking agents, colouring agents and corrosion inhibitors. These adjuvants may also serve as carriers or diluents.

Granular formulations may be prepared by absorbing the compounds of the present invention (dissolved in suitable solvents, which may, if desired, be volatile) onto the solid diluents or carriers in granular form and, if desired, evaporating the solvents, or by granulating compositions in powder form obtained as described above.

Powders, dispersible granules and liquid compositions in the form of concentrates may be diluted with water or other suitable diluents, for example mineral or vegetable oils, particularly in the case of liquid concentrates in which the diluent or carrier is an oil, to give compositions ready for use.

The wettable powders (or powders for spraying) usually contain from 20 to 95% of combination, and they usually contain, in addition to the solid vehicle, from 0 to 5% of a wetting agent, from 3 to 10% of a dispersant agent and if necessary, from 0 to 10% of one or more stabilisers and/or other additives such as penetrating agents, adhesives or anti-caking agents and colourings.

The aqueous suspension concentrates, which are applicable by spraying, are prepared in such a way as to obtain a stable fluid product (by fine grinding) which does not settle out and they usually contain from 10 to 75% of combination, from 0.5 to 15% of surface acting agents, from 0.1 to 10% of thixotropic agents, from 0 to 10% of suitable

additives such as antifoams, corrosion inhibitors, stabilisers, and water or an organic liquid in which the active substance is sparingly soluble or insoluble. Some organic solid substances or inorganic salts can be dissolved in order to assist in preventing sedimentation or as antifreeze for the water.

Application of the herbicide, antidote, or mixture thereof, can be carried out by conventional techniques utilizing, for example, hand-carried or tractor-mounted spreaders, power dusters, boom and hand sprayers, spray dusters, and granular applicators. If desired, application of the compositions of the invention to plants can be accomplished by incorporating the compositions in the soil or other media.

Herbicidal compositions according to the present invention may also comprise (a) and (b) in association with, and preferably homogeneously dispersed in, one or more other pesticidally active compounds and, if desired one or more compatible pesticidally acceptable diluents and carriers. Examples of other pesticidally active ingredients include fungicides, insecticides, plant growth regulators and, most preferably, herbicides.

The optional partner herbicides which may be combined with the derivatives of formula (I) and antidote are preferably selected from chloroacetamides (e.g. metolachlor, acetochlor, alachlor), sulfonyleureas, thiocarbamates, dithiocarbamates, metribuzin, sulfentrazone, flumetsulam, clorasulam-methyl, oxasulfuron, flumiclorac, bentazon, chlorimuron, linuron, clomazone, dimethenamid, pendimethalin, trifluralin, clethodim and acifluorfen, bifenox, diflufenican, diuron, atrazine and ametryne.

According to a further feature of the present invention there is provided a product comprising:

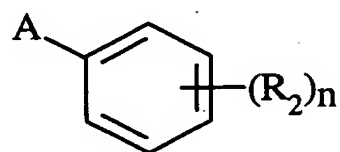
(a) a herbicidally effective amount of a benzoylisoxazole and/or dione derivative of formula (I), or an agriculturally acceptable salt or metal complex thereof; and

(b) an antidotally effective amount of an antidote compound or an agriculturally acceptable salt thereof;

as a combined preparation for separate, simultaneous or sequential use in the control of weeds at a crop locus.

CLAIMS

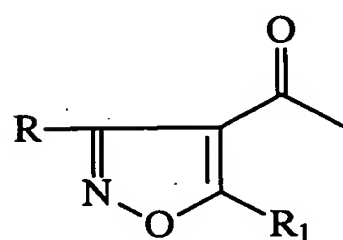
1. A method of reducing phytotoxicity to a crop (especially maize)
at a locus caused by the application thereto of a herbicidal
benzoylisoxazole and/or dione derivative of formula (I):



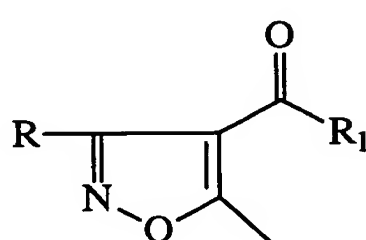
(I)

wherein:

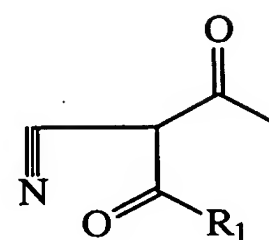
A represents a group (A-1) to (A-7):



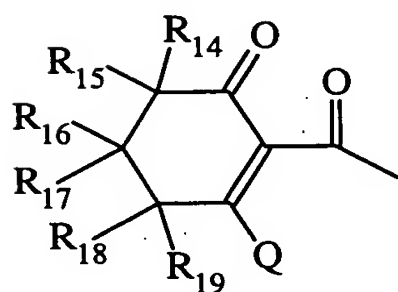
(A-1)



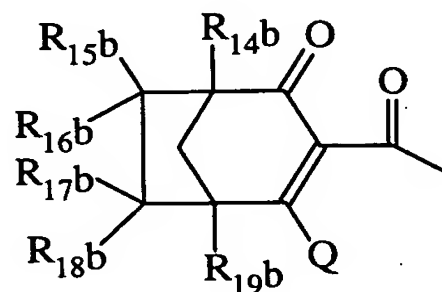
(A-2)



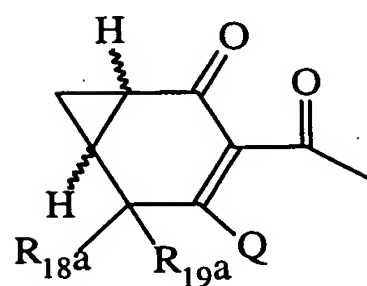
(A-3)



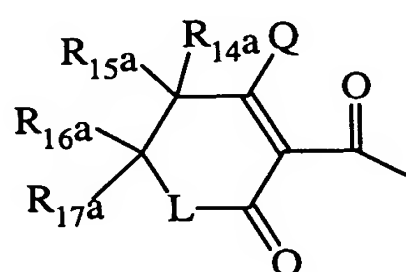
(A-4)



(A-5)

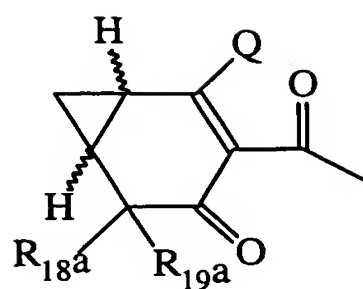


(A-6)

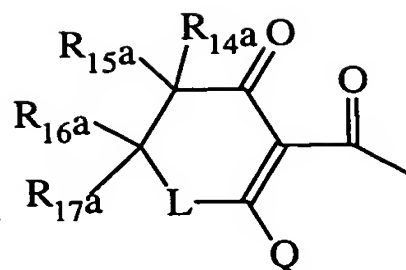


(A-7)

or a corresponding formula (A-6a) or (A-7a):



(A-6a)



(A-7a)

in which the position of the carbonyl group and the group Q are reversed and the double bond in the ring is attached to the carbon atom attached to the group Q;

R represents a hydrogen atom or a halogen atom; a straight- or branched chain alkyl, alkenyl or alkynyl group containing from one to six carbon atoms which is optionally substituted by one or more halogen atoms; a cycloalkyl group containing from 3 to 6 carbon atoms optionally substituted by one or more groups R^5 , one or more halogen atoms or a group $-CO_2R^3$; or a group selected from $-CO_2R^3$, $-COR^5$, cyano, nitro, $-CONR^3R^4$ and $-S(O)_kR^{13}$;

R^1 represents a straight- or branched-chain alkyl, alkenyl or alkynyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms; or a cycloalkyl group containing from three to six carbon atoms optionally substituted by one or more groups R^5 or one or more halogen atoms;

R^2 represents a halogen atom; a straight- or branched-chain alkyl, alkenyl or alkynyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms; a straight- or branched-chain alkyl group containing up to six carbon atoms which is substituted by one or more groups $-OR^5$; or a group selected from nitro, cyano, $-CO_2R^5$, $-S(O)_pR^6$, $-O(CH_2)_mOR^5$, $-COR^5$, $-NR^{11}R^{12}$, $-N(R^8)SO_2R^7$, $-N(R^8)CO_2R^7$, $-OR^5$, $-OSO_2R^7$, $-SO_2NR^3R^4$, $-CONR^3R^4$, $-CSNR^3R^4$, $-(CR^9R^{10})_t-S(O)_qR^7$ and $-SF_5$;

or two groups R^2 , on adjacent carbon atoms of the phenyl ring may, together with the carbon atoms to which they are attached, form a 5 to 7 membered saturated or unsaturated heterocyclic ring containing up to three ring heteroatoms selected from nitrogen, oxygen and sulfur, which ring is optionally substituted by one or more groups selected from halogen, nitro, $-S(O)_pR^{13}$, C_{1-4} alkyl, C_{1-4} alkoxy, C_{1-4} haloalkyl, C_{1-4} haloalkoxy, $=O$ (or a 5- or 6- membered cyclic acetal thereof), and $=NO-R^3$, it being understood that a sulphur atom, where present in the ring, may be in the form of a group $-SO-$ or $-SO_2-$;

n represents an integer from one to five: when n is greater than one the groups R^2 may be the same or different;

R^3 , R^4 and R^{22} each independently represent a hydrogen atom, or a straight- or branched chain alkyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms;

R^5 and R^{23} each independently represent a straight- or branched-chain alkyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms or a straight- or branched-chain alkenyl or alkynyl group containing from two to six (preferably from three to six) carbon atoms which is optionally substituted by one or more halogen atoms;

R^6 and R^7 , which may be the same or different, each represent R^5 ; or phenyl optionally substituted by from one to five groups which may be the same or different selected from a halogen atom, a straight- or branched-chain alkyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms, nitro, cyano, $-CO_2R^5$, $-S(O)_pR^{13}$, $-NR^{11}NR^{12}$, $-OR^5$ and $-CONR^3R^4$;

R^8 , R^9 and R^{10} each represent a hydrogen atom or R^6 ;

R^{11} and R^{12} each represent hydrogen or R^5 ;

R^{13} and R^{21} represent a straight- or branched-chain alkyl group containing up to six carbon atoms which is optionally substituted by one or more halogen atoms;

Q represents hydroxy, C1-6 alkoxy, OR^{20} , SR^{20} or SR^{21} ;

5 L represents oxygen or NR^{22} ;

R^{14} , R^{14a} , R^{14b} , R^{15} , R^{15a} , R^{15b} , R^{16} , R^{16a} , R^{16b} , R^{17} , R^{17a} , R^{17b} , R^{18} , R^{18a} , R^{18b} , R^{19} , R^{19a} and R^{19b} represent the same or different groups selected from hydrogen, R^{23} , $-(CH_2)_uCO_2R^{22}$, halogen, cyano, C1-6 alkoxy, $-(CH_2)_x$ -[phenyl optionally substituted by
10 from one to five groups R^{24} which may be the same or different], and cycloalkyl containing from three to six carbon atoms optionally substituted by C1-6 alkyl or $-S(O)_pR^{21}$;

R^{20} represents phenyl optionally substituted by from one to five groups selected from halogen, C1-6 alkyl, C1-6 haloalkyl, C1-6 alkoxy
15 and nitro;

R^{24} represents a group selected from halogen, R^{25} , nitro, cyano, $-CO_2R^{26}$, $-S(O)_pR^{21}$, $-OR^{21}$ and $-NR^{26}R^{27}$;

R^{25} represents a straight- or branched- chain alkyl group containing one to three carbon atoms optionally substituted by one or
20 more halogen atoms;

R^{26} and R^{27} which may be the same or different, each represents hydrogen or R^{23} ;

p, q and u independently represent the values zero, one or two;

k and m represent one, two or three;

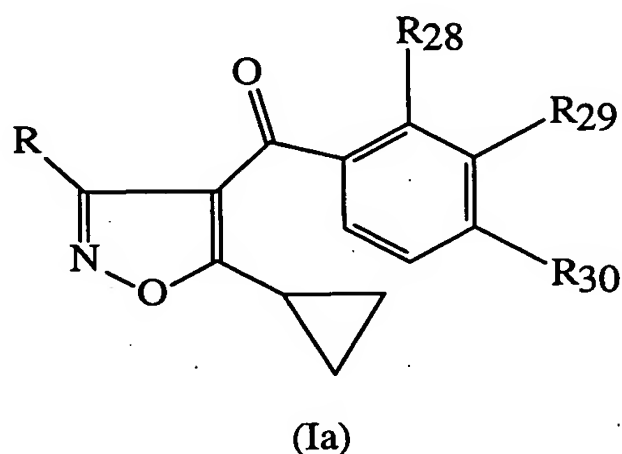
25 x represents zero or one;

t represents an integer from one to four; when t is greater than one, the groups R^9 and R^{10} may be the same or different;

or an agriculturally acceptable salt or metal complex thereof;
which method comprises applying to the locus of the crop an antidotally

effective amount of an antidote compound, optionally with a partner herbicide.

2. A method according to claim 1 in which the isoxazole or dione herbicide has the general formula (Ia):



wherein:

R is hydrogen or $-\text{CO}_2\text{Et}$;

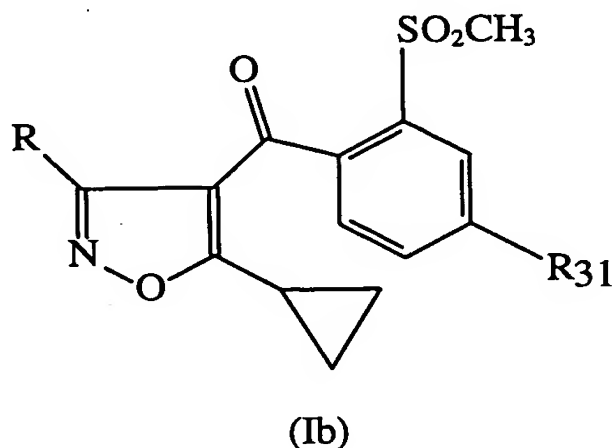
R^{28} is selected from $-\text{S}(\text{O})_p\text{Me}$, Me, Et, a chlorine, bromine or fluorine atom, methoxy, ethoxy and $-\text{CH}_2\text{S}(\text{O})_q\text{Me}$;

R^{29} is selected from a hydrogen atom, a chlorine, bromine or fluorine atom, methoxy, ethoxy and $-\text{S}(\text{O})_p\text{Me}$;

R^{30} is selected from a hydrogen atom, a chlorine, bromine or fluorine atom, methoxy and trifluoromethyl; and

p and q independently represent the values zero, one or two.

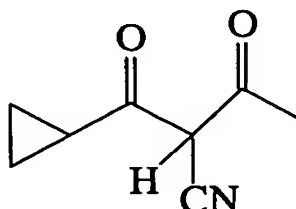
3. A method according to claim 1 or 2 in which the isoxazole or dione herbicide has the general formula (Ib):



wherein R^{31} is chlorine, bromine or trifluoromethyl; and

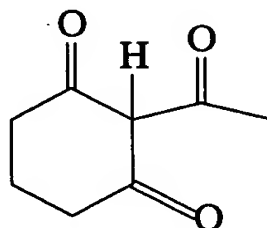
R is hydrogen or $-\text{CO}_2\text{Et}$.

4. A method according to any one of the preceding claims in which a substituted phenyl ring as defined in formulae (I); (Ia); or (Ib) as depicted in claim 11,12 or 13 is attached to a grouping;



5. A method according to claim 4 in which the phenyl ring is substituted by two groups independently selected from halogen, alkyl, S(O)_talkyl (t = 0, 1 or 2) or haloalkyl.

6. A method according to claim 1, 2 or 3 in which a substituted phenyl ring as defined above in formula (I); (Ia); or (Ib) is attached to a grouping;



7. A method according to claim 1 wherein the compound of formula (I) is:

5-cyclopropyl-4-[2-chloro-3-ethoxy-4-(ethylsulphonyl)benzoyl]isoxazole;

4-(4-chloro-2-methylsulphonylbenzoyl)-5-cyclopropylisoxazole;

5-cyclopropyl-4-(2-methylsulphonyl-4-trifluoromethylbenzoyl)isoxazole;

4-(4-bromo-2-methylsulphonylbenzoyl)-5-cyclopropylisoxazole;

5-cyclopropyl-4-[4-fluoro-3-methoxy-2-(methylsulphonyl)benzoyl]isoxazole;

4-(4-bromo-2-methylsulphonylmethylbenzoyl)-5-cyclopropylisoxazole;

ethyl 5-cyclopropyl-4-(2-methylsulphonyl-4-trifluoromethylbenzoyl) isoxazole-3-carboxylate;

2-[2-chloro-(4-methylsulphonyl)benzoyl]-1,3-cyclohexanedione;

2-[2-nitro-(4-methylsulphonyl)benzoyl]-1,3-cyclohexanedione;
2-(2,3-dihydro-5,8-dimethyl-1,1-dioxospiro[4H-1-benzothiine-4,2'
[1,3]dioxolan]-6-ylcarbonyl)cyclohexane-1,3-dione;
5-cyclopropyl-4-(2-methylsulphonyl-4-trifluoromethylbenzoyl)-3-
5 methylthio-isoxazole;
2-cyano-3-cyclopropyl-1-(2-methylsulphonyl-4-
trifluoromethylphenyl) propan-1,3-dione.

8. A method according to claim 17 in which the compound is
5-cyclopropyl-4-(2-methylsulphonyl-4-
10 trifluoromethylbenzoyl)isoxazole or 2-[2-nitro-(4-
methylsulphonyl)benzoyl]-1,3-cyclohexanedione.

9. A method according to claim 1 in which the antidote is
selected from:

flurazole; fenclorazole-ethyl; fenchorazole; benoxacor;
15 dichlormid; fenclorim; furilazole; mefenpyr-diethyl; CMPI; 4-hydroxy-
1-methyl-3-(1-1*H*-tetrazol-5-ylmethanoyl)-1*H*-quinolin-2-one;
daimuron; (S)-MBU; dimepiperate; 5,5-diphenylisoxazolinone-3-
carboxylic acid; and ethyl 5,5-diphenylisoxazolinone-3-carboxylate.

10. A method according to any one of the preceding claims in
20 which the crop plant to be protected is maize.

11. A method according to claim 1 in which the application
rate of the benzoylisoxazole and/or dione of formula (I) is from 0.004kg
to 5kg per hectare.

12. A method according to claim 1 in which the application
25 rate of the benzoylisoxazole and/or dione of formula (I) is from 0.01kg
to 2kg per hectare.

13. A herbicidal composition comprising:

(a) a herbicidally effective amount of a benzoylisoxazole
and/or dione derivative of formula (I) or an agriculturally acceptable salt
30 or metal complex thereof, optionally in combination with a partner
herbicide; and

(b) an antidotally effective amount of an antidote compound;

in association with a herbicidally acceptable diluent or carrier and/or surface active agent.

14. A composition according to claim 13 in which the weight ratio of the compound of formula (I):antidote is from 1:25 to 60:1.

5

15. A product comprising:

(a) a herbicidally effective amount of a benzoylisoxazole and/or dione derivative of formula (I), or an agriculturally acceptable salt or metal complex thereof; and

10

(b) an antidotally effective amount of an antidote; wherein said antidote is antidotally effective to said benzoylisoxazole and/or dione derivative;

as a combined preparation for separate, simultaneous or sequential use in the control of weeds at a locus.

15

16. A method according to claim 1 substantially as hereinbefore described.

17. A composition according to claim 13 substantially as hereinbefore described.

20

18. A product according to claim 15 substantially as hereinbefore described.

ABSTRACT

The present invention provides a method of reducing phytotoxicity to crops (especially maize) caused by a herbicidal benzoylisoxazole and/or dione derivative of formula (I) or an agriculturally acceptable salt or metal complex thereof; which method comprises applying to the locus of the crop an antidotally effective amount of an antidote compound, optionally with a partner herbicide.

